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Inventors: Ronald S. Cok
Customer No. 01333

OLED DISPLAY WITH CIRCULAR POLARIZER

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P.O. Box 1450
Alexandria, VA. 22313-1450

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OLED DISPLAY WITH CIRCULAR POLARIZER

CROSS-REFERENCE TO RELATED APPLICATIONS

This is a continuation-in-part of copending application USSN
5 10/271,149, filed October 15, 2002, the disclosure of which is incorporated by
reference in its entirety.

FIELD OF THE INVENTION

The present invention relates to organic light emitting diode
10 (OLED) displays and, more particularly, to such displays having circular
polarizing elements to reduce glare and increase the display contrast.

BACKGROUND OF THE INVENTION

Emissive flat-panel display devices are widely used in conjunction
15 with computing devices and in particular with portable devices. These displays
are often used in portable devices or in public areas with significant ambient
illumination. In these locations, the contrast of the display is of great concern.

In particular, OLED display devices suffer from problems with
contrast. It is known to use a circular light polarizer affixed to the surface of the
20 display so that light incident on the display is absorbed by the polarizer, while
light emitted by the display is not. This is problematic in that the circular light
polarizer is exposed to the environment and is subject to scratching, peeling,
moisture, dents, and the like, which reduces its effectiveness and acceptability.

In an attempt to address the problem, WO0210845 A2 entitled
25 "High Durability Circular Polarizer for use with Emissive Displays" published
February 7, 2002 describes a high durability circular light polarizer including an
unprotected K-type polarizer and a quarter-wavelength retarder and designed for
use with an emissive display module such as an organic light emitting diode or a
plasma display device. Such devices are expensive and remain subject to
30 environmental stress which can degrade their performance. Moreover, placing a
circular light polarizer on the surface of the display device inhibits the further

integration of other elements such as lenslet arrays and touch screen components over the display.

An additional protective cover may be provided over the circular light polarizer. The LS633 digital camera, sold by Eastman Kodak Company, e.g., employed a polycarbonate cover located over the circular light polarizer to provide environmental protection to the OLED display and circular light polarizer. While such an approach is effective in protecting the display and circular light polarizer, it adds weight, size, and cost to the display. Moreover, modeling and tests done by applicant demonstrate that such a design severely reduces the contrast of the display.

There is a need therefore for an improved OLED display that improves the robustness of the display while maintaining the display contrast.

SUMMARY OF THE INVENTION

The need is met according to the present invention by providing a top-emitting OLED display that includes a substrate; an array of OLED light emissive elements formed over the substrate; an encapsulating cover located over the OLED light emissive elements; and a circular light polarizer located between the encapsulating cover and the OLED light emissive elements.

ADVANTAGES

The present invention has the advantage that it improves the contrast and robustness of an OLED display by protecting the circular light polarizer from environmental wear and enables the application of additional structures on the top of the encapsulating cover.

BRIEF DESCRIPTION OF THE DRAWINGS

Fig. 1 is a schematic diagram of a prior-art OLED;

Fig. 2 is a partial cross section of a prior-art top-emitting OLED display device;

Fig. 3 is a cross section of a top-emitting OLED display device with a circular light polarizer located on the top of the display as suggested by the prior art;

Fig. 4 is a cross section of a top-emitting OLED display according to one embodiment of the present invention;

Fig. 5 is a cross section of a top-emitting OLED display according to an alternative embodiment of the present invention;

Fig. 6 is a cross section of a top-emitting OLED display according to yet another alternative embodiment of the present invention;

Fig. 7 is a cross section of a top-emitting OLED display according to yet another alternative embodiment of the present invention;

Fig. 8 is a cross section of a top-emitting OLED display according to yet another alternative embodiment of the present invention; and

Fig. 9 is a partial cross section of a prior art OLED emitter having multiple layers.

DETAILED DESCRIPTION OF THE INVENTION

Referring to Fig. 1, a prior-art OLED includes a substrate **20** such as glass or plastic and an OLED light emissive element **10** having an organic light-emitting layer **12** disposed between two electrodes, e.g. a cathode **14** and an anode **16**. The organic light emitting layer **12** emits light upon application of a voltage from a power source **18** across the electrodes. It will be understood that the relative locations of the electrodes **14** and **16** may be reversed with respect to the substrate. The light-emitting layer **12** may include other layers such as electron-injection or hole-injection layers as is known in the art.

Referring to Fig. 2, a prior-art top-emitting OLED display device **11** includes a substrate **20**, a thin-film transistor (TFT) active-matrix layer **22** that provides power to an OLED light emitting layer **12**. A patterned first planarizing insulating layer **24** is provided over the TFT active-matrix layer, and an array of first electrodes **16** are provided over the planarized insulating layer **24** and in electrical contact with the TFT active matrix layer. A patterned second insulating layer **24'** is provided over the array of first electrodes **16** such that at least a

portion of the each of the first electrodes **16** is exposed and the various electrodes **16** do not form an electrical short circuit.

Over the first electrodes and insulating layers are provided red, green, and blue-emitting organic OLED elements, **12R**, **12G**, and **12B**,
5 respectively. These elements are composed of further layers as described in more detail below. Herein, the collection of OLED elements, including hole-injection **26**, hole-transport **27**, electron-injection **29**, and electron-transport layers **28**, may also be referred to as the OLED light-emitting layer **12**. The light-emitting area is generally defined by the area of the first electrode **16** in contact with the OLED
10 elements. Over the OLED light-emitting layer **12** is provided a transparent, common second electrode **14** that has sufficient optical transparency to allow transmission of the generated red, green, and blue light. An optional second electrode protection layer **32** may be provided to protect the electrode and underlying layers. Each first electrode in combination with its associated OLED
15 element and second electrode is herein referred to as an OLED light emissive element **10**. A typical top-emitting OLED display device comprises an array of OLED light-emitting elements wherein each OLED light-emitting elements emits red, green or blue light. A cavity **34** generally filled with inert gas or a transmissive polymer material, separates the optional electrode protection layer
20 from an encapsulating cover **36**.

Referring to Fig. 3, a prior-art top-emitting OLED may be provided with a circular light polarizer **50** that has the property that it will polarize light that passes through the polarizer and will absorb such polarized light that is reflected from the OLED light emissive elements **10** or substrate **20**. About half of the light
25 emitted from the light emissive elements **10** passes through the circular light polarizer, but most of the ambient light falling on the device is absorbed by the circular light polarizer. As noted above, the problem with this arrangement is that the circular light polarizer is subjected to the external environment and can be easily scratched and is subject to delamination from the surface of the display
30 device. An optional protective cover **38** may be provided but, as described above, such a cover adds weight, thickness, and cost to the display and reduces contrast by introducing new reflective interfaces.

Referring to Fig. 4, according to the present invention, the circular light polarizer 50 is located between the encapsulating cover and the OLED light emissive elements, thereby protecting the circular light polarizer from the environment. In the preferred embodiment, the encapsulating cover 36 defines a cavity 34 and is affixed to the substrate 20 by a suitable adhesive 70, typically an epoxy. The cavity 34 may be sufficiently deep to leave a gap between the circular light polarizer 50 and the OLED light emissive elements 10. The present invention may include the optional electrode protection layer 32 shown in Fig. 2 to further protect the electrode 14 and layers beneath the electrode. Moreover, the adhesive 70, if it is sufficiently transparent may be used to fill the cavity 34 between the circular light polarizer and the OLED light emissive elements 10. The circular light polarizer 50 may be attached to the inside of the encapsulating cover 36 with a suitable adhesive.

Circular light polarizers are typically made with layers of polymeric materials. In a preferred embodiment, the material used to fill cavity 34 has a refractive index matched more closely than air to the refractive index of the adjacent surface layer of the circular light polarizer, preferably matched to at most 10% difference, more preferably matched to at most 5% difference, and most preferably matched to at most 1% difference. Use of materials in adjacent layers having a common refractive index will reduce reflections and optical interference. For example, the linear polarizer layers of a circular light polarizer can be made with polymers such as triacetate cellulose which has a refractive index of 1.487. More generally, polymers may have a refractive index generally between 1.45 and 1.6 and may be used as a material to fill cavity 34. Theoretically, a material used to fill the cavity 34 may alternatively have a refractive index matched to that of the cathode or cathode protective layer. However, materials conventionally used for a cathode (such as silver or indium tin oxide) have a refractive index very different from that of glass or plastic. For example, indium tin oxide has a refractive index of approximately 1.9.

An adhesive may also be applied between the circular light polarizer 50 and the encapsulating cover 36. If such an adhesive is used, it is preferred that the refractive index of the adhesive also match the refractive index

of the adjacent surface of the circular light polarizer. Adhesives with such refractive indices are known. Alternatively, such an adhesive may have a refractive index matched to the encapsulating cover 36. Such covers have a refractive index in the region of 1.5, for example borosilicate glass having a refractive index of 1.51.

In a preferred embodiment, an anti-reflection coating 37 is provided on the encapsulating cover 36 on the side opposed from the circular light polarizer. Such coatings are known in the art and further reduce reflections and optical interference in the display. Additional coatings, such as hardeners, may also be provided to provide environmental protection to the encapsulating cover 36.

Referring to Fig. 5, in an alternative embodiment of the present invention, the circular light polarizer 50 is located on top of the OLED light emissive elements 10. In this arrangement, any adhesive used to adhere the circular light polarizer 50 to the OLED light emissive elements 10 should have a refractive index preferably matched to the adjacent surface layer of the circular light polarizer 50 or, alternatively, to the top layer of the OLED light emissive elements 10.

Referring to Fig. 6, in a still further alternative embodiment of the present invention, the encapsulating cover 36 does not define a cavity. The circular light polarizer 50 is attached to the encapsulating cover 36. A transparent adhesive layer 70 hermetically seals the perimeter of the encapsulating cover over the OLED light emissive elements 10 and may extend over the OLED light emissive elements 10. Alternatively, the circular light polarizer 50 may be attached to the OLED light emissive elements 10 and the transparent adhesive extends between the circular light polarizer and the encapsulating cover. In either, case, it is preferred that the adhesive layer 70 have a refractive index matched to that of the adjacent surface layer of the circular light polarizer 50. According to a further alternative, the adhesive 70 may be located only around the periphery of the encapsulating cover and can comprise a light absorbing material.

Referring to Figs. 7 and 8, the encapsulating cover 36 may be provided with a peripheral channel 52 that is filled with a desiccant material. Fig.

7 shows such an arrangement wherein the encapsulating cover defines a cavity over the OLED light emissive elements, and Fig. 8 shows the arrangement wherein the encapsulating cover does not. In either case, a gap filled with an inert gas or light transmissive material may be provided between the circular light polarizer and the encapsulating cover or the OLED light emissive elements. The circular light polarizer may be affixed to the OLED light emissive elements 10 as shown in Fig. 5. As described above, if any cavity filling material is used and located in contact with the circular light polarizer 50, it is preferred that the material have a refractive index matched to that of the adjacent surface layer of the circular light polarizer.

In all of the cases described in Figs. 4, 5, 6, 7, and 8, it is preferred that an anti-reflective coating be used on the outside face of the encapsulating cover in conjunction with optional environmentally protective coatings. If the cavity 34 is filled with a material, the material preferably has a refractive index matched to that of the adjacent surface layer of the circular light polarizer. Moreover, if the cavity 34 is filled with material, an additional adhesive between the circular light polarizer 50 and the encapsulating cover 36 may not be necessary as the filler material may hold the circular light polarizer in place. Likewise, if the circular light polarizer 50 is located on the OLED light emissive element 10 (as in Fig. 5), an additional adhesive may not be necessary.

In a preferred embodiment, the invention is employed in a device that includes Organic Light Emitting Diodes (OLEDs) which are composed of small molecule or polymeric OLEDs as disclosed in but not limited to US 4,769,292, issued September 6, 1988 to Tang et al. and US 5,061,569, issued October 29, 1991 to VanSlyke et al. Many combinations and variations of organic light emitting displays can be used to fabricate such a device.

General device architecture

The present invention can be employed in most OLED material configurations. These include very simple structures comprising a single anode and cathode to more complex devices, such as passive matrix displays comprised of orthogonal arrays of anodes and cathodes to form pixels, and active-matrix

displays where each pixel is controlled independently, for example, with thin film transistors (TFTs).

There are numerous configurations of the organic layers wherein the present invention can be successfully practiced. A typical structure is shown in Fig. 9 and is comprised of a substrate **101**, an anode **103**, a hole-injecting layer **105**, a hole-transporting layer **107**, a light-emitting layer **109**, an electron-transporting layer **111**, and a cathode **113**. These layers are described in detail below. Note that the substrate may alternatively be located adjacent to the cathode, or the substrate may actually constitute the anode or cathode. The organic layers between the anode and cathode are conveniently referred to as the organic EL element. The total combined thickness of the organic layers is preferably less than 500 nm.

The anode and cathode of the OLED are connected to a voltage/current source **250** through electrical conductors **260**. The OLED is operated by applying a potential between the anode and cathode such that the anode is at a more positive potential than the cathode. Holes are injected into the organic EL element from the anode and electrons are injected into the organic EL element at the cathode. Enhanced device stability can sometimes be achieved when the OLED is operated in an AC mode where, for some time period in the cycle, the potential bias is reversed and no current flows. An example of an AC driven OLED is described in US 5,552,678.

Substrate

The OLED device of this invention is typically provided over a supporting substrate where either the cathode or anode can be in contact with the substrate. The electrode in contact with the substrate is conveniently referred to as the bottom electrode. Conventionally, the bottom electrode is the anode, but this invention is not limited to that configuration. The substrate can either be transmissive or opaque. In the case wherein the substrate is transmissive, a reflective or light absorbing layer is used to reflect the light through the encapsulating cover or to absorb the light, thereby improving the contrast of the display. Substrates can include, but are not limited to, glass, plastic,

semiconductor materials, silicon, ceramics, and circuit board materials. Of course it is necessary to provide a light-transparent top electrode.

Anode

When EL emission is viewed through anode **103**, the anode should
5 be transparent or substantially transparent to the emission of interest. Common transparent anode materials used in this invention are indium-tin oxide (ITO), indium-zinc oxide (IZO) and tin oxide, but other metal oxides can work including, but not limited to, aluminum- or indium-doped zinc oxide, magnesium-indium oxide, and nickel-tungsten oxide. In addition to these oxides, metal nitrides, such
10 as gallium nitride, and metal selenides, such as zinc selenide, and metal sulfides, such as zinc sulfide, can be used as the anode. For applications where EL emission is viewed only through the cathode electrode, the transmissive characteristics of anode are immaterial and any conductive material can be used, transparent, opaque or reflective. Example conductors for this application
15 include, but are not limited to, gold, iridium, molybdenum, palladium, and platinum. Typical anode materials, transmissive or otherwise, have a work function of 4.1 eV or greater. Desired anode materials are commonly deposited by any suitable means such as evaporation, sputtering, chemical vapor deposition, or electrochemical means. Anodes can be patterned using well-known
20 photolithographic processes. Optionally, anodes may be polished prior to application of other layers to reduce surface roughness so as to minimize shorts or enhance reflectivity.

Hole-Injecting Layer (HIL)

While not always necessary, it is often useful to provide a hole-
25 injecting layer **105** between anode **103** and hole-transporting layer **107**. The hole-injecting material can serve to improve the film formation property of subsequent organic layers and to facilitate injection of holes into the hole-transporting layer. Suitable materials for use in the hole-injecting layer include, but are not limited to, porphyrinic compounds as described in US 4,720,432, plasma-deposited
30 fluorocarbon polymers as described in US 6,208,075, and some aromatic amines, for example, m-MTDATA (4,4',4''-tris[(3-methylphenyl)phenylamino]triphenylamine). Alternative hole-injecting materials

reportedly useful in organic EL devices are described in EP 0 891 121 A1 and EP 1 029 909 A1.

Hole-Transporting Layer (HTL)

The hole-transporting layer **107** contains at least one hole-transporting compound such as an aromatic tertiary amine, where the latter is understood to be a compound containing at least one trivalent nitrogen atom that is bonded only to carbon atoms, at least one of which is a member of an aromatic ring. In one form the aromatic tertiary amine can be an arylamine, such as a monoarylamine, diarylamine, triarylamine, or a polymeric arylamine. Exemplary monomeric triarylaminines are illustrated by Klupfel et al. in US 3,180,730. Other suitable triarylaminines substituted with one or more vinyl radicals and/or comprising at least one active hydrogen containing group are disclosed by Brantley et al. in US 3,567,450 and 3,658,520.

A more preferred class of aromatic tertiary amines are those which include at least two aromatic tertiary amine moieties as described in US 4,720,432 and 5,061,569. The hole-transporting layer can be formed of a single or a mixture of aromatic tertiary amine compounds. Illustrative of useful aromatic tertiary amines are the following:

1,1-Bis(4-di-*p*-tolylaminophenyl)cyclohexane
1,1-Bis(4-di-*p*-tolylaminophenyl)-4-phenylcyclohexane
4,4'-Bis(diphenylamino)quadriphenyl
Bis(4-dimethylamino-2-methylphenyl)-phenylmethane
N,N,N-Tri(*p*-tolyl)amine
4-(di-*p*-tolylamino)-4'-[4(di-*p*-tolylamino)-styryl]stilbene
N,N,N',N'-Tetra-*p*-tolyl-4-4'-diaminobiphenyl
N,N,N',N'-Tetraphenyl-4,4'-diaminobiphenyl
N,N,N',N'-tetra-1-naphthyl-4,4'-diaminobiphenyl
N,N,N',N'-tetra-2-naphthyl-4,4'-diaminobiphenyl
N-Phenylcarbazole
4,4'-Bis[N-(1-naphthyl)-N-phenylamino]biphenyl
4,4'-Bis[N-(1-naphthyl)-N-(2-naphthyl)amino]biphenyl
4,4"-Bis[N-(1-naphthyl)-N-phenylamino]p-terphenyl

4,4'-Bis[N-(2-naphthyl)-N-phenylamino]biphenyl
 4,4'-Bis[N-(3-acenaphthenyl)-N-phenylamino]biphenyl
 1,5-Bis[N-(1-naphthyl)-N-phenylamino]naphthalene
 4,4'-Bis[N-(9-anthryl)-N-phenylamino]biphenyl
 5 4,4''-Bis[N-(1-anthryl)-N-phenylamino]-*p*-terphenyl
 4,4'-Bis[N-(2-phenanthryl)-N-phenylamino]biphenyl
 4,4'-Bis[N-(8-fluoranthryl)-N-phenylamino]biphenyl
 4,4'-Bis[N-(2-pyrenyl)-N-phenylamino]biphenyl
 4,4'-Bis[N-(2-naphthacenyl)-N-phenylamino]biphenyl
 10 4,4'-Bis[N-(2-perylenyl)-N-phenylamino]biphenyl
 4,4'-Bis[N-(1-coronenyl)-N-phenylamino]biphenyl
 2,6-Bis(di-*p*-tolylamino)naphthalene
 2,6-Bis[di-(1-naphthyl)amino]naphthalene
 2,6-Bis[N-(1-naphthyl)-N-(2-naphthyl)amino]naphthalene
 15 N,N,N',N'-Tetra(2-naphthyl)-4,4''-diamino-*p*-terphenyl
 4,4'-Bis{N-phenyl-N-[4-(1-naphthyl)-phenyl]amino}biphenyl
 4,4'-Bis[N-phenyl-N-(2-pyrenyl)amino]biphenyl
 2,6-Bis[N,N-di(2-naphthyl)amine]fluorene
 1,5-Bis[N-(1-naphthyl)-N-phenylamino]naphthalene
 20 4,4',4''-tris[(3-methylphenyl)phenylamino]triphenylamine

Another class of useful hole-transporting materials includes polycyclic aromatic compounds as described in EP 1 009 041. Tertiary aromatic amines with more than two amine groups may be used including oligomeric materials. In addition, polymeric hole-transporting materials can be used such as
 25 poly(N-vinylcarbazole) (PVK), polythiophenes, polypyrrole, polyaniline, and copolymers such as poly(3,4-ethylenedioxythiophene) / poly(4-styrenesulfonate) also called PEDOT/PSS.

Light-Emitting Layer (LEL)

As more fully described in US 4,769,292 and 5,935,721, the light-emitting layer (LEL) 109 of the organic EL element includes a luminescent or
 30 fluorescent material where electroluminescence is produced as a result of electron-hole pair recombination in this region. The light-emitting layer can be comprised

of a single material, but more commonly consists of a host material doped with a guest compound or compounds where light emission comes primarily from the dopant and can be of any color. The host materials in the light-emitting layer can be an electron-transporting material, as defined below, a hole-transporting material, as defined above, or another material or combination of materials that support hole-electron recombination. The dopant is usually chosen from highly fluorescent dyes, but phosphorescent compounds, e.g., transition metal complexes as described in WO 98/55561, WO 00/18851, WO 00/57676, and WO 00/70655 are also useful. Dopants are typically coated as 0.01 to 10 % by weight into the host material. Polymeric materials such as polyfluorenes and polyvinylarylenes (e.g., poly(p-phenylenevinylene), PPV) can also be used as the host material. In this case, small molecule dopants can be molecularly dispersed into the polymeric host, or the dopant could be added by copolymerizing a minor constituent into the host polymer.

15 An important relationship for choosing a dye as a dopant is a comparison of the bandgap potential which is defined as the energy difference between the highest occupied molecular orbital and the lowest unoccupied molecular orbital of the molecule. For efficient energy transfer from the host to the dopant molecule, a necessary condition is that the band gap of the dopant is smaller than that of the host material. For phosphorescent emitters it is also important that the host triplet energy level of the host be high enough to enable energy transfer from host to dopant.

Host and emitting molecules known to be of use include, but are not limited to, those disclosed in US 4,768,292; 5,141,671; 5,150,006; 5,151,629; 25 5,405,709; 5,484,922; 5,593,788; 5,645,948; 5,683,823; 5,755,999; 5,928,802; 5,935,720; 5,935,721; and 6,020,078.

Metal complexes of 8-hydroxyquinoline (oxine) and similar derivatives constitute one class of useful host compounds capable of supporting electroluminescence. Illustrative of useful chelated oxinoid compounds are the following:

CO-1: Aluminum trisoxine [alias, tris(8-quinolinolato)aluminum(III)]

CO-2: Magnesium bisoxine [alias, bis(8-quinolinolato)magnesium(II)]

CO-3: Bis[benzo{f}-8-quinolinolato]zinc (II)

CO-4: Bis(2-methyl-8-quinolinolato)aluminum(III)- μ -oxo-bis(2-methyl-8-quinolinolato) aluminum(III)

CO-5: Indium trisoxine [alias, tris(8-quinolinolato)indium]

5 CO-6: Aluminum tris(5-methyloxine) [alias, tris(5-methyl-8-quinolinolato)aluminum(III)]

CO-7: Lithium oxine [alias, (8-quinolinolato)lithium(I)]

CO-8: Gallium oxine [alias, tris(8-quinolinolato)gallium(III)]

CO-9: Zirconium oxine [alias, tetra(8-quinolinolato)zirconium(IV)]

10 Other classes of useful host materials include, but are not limited to: derivatives of anthracene, such as 9,10-di-(2-naphthyl)anthracene and derivatives thereof as described in US 5,935,721, distyrylarylene derivatives as described in US 5,121,029, and benzazole derivatives, for example, 2, 2', 2''-(1,3,5-phenylene)tris[1-phenyl-1H-benzimidazole]. Carbazole derivatives are
15 particularly useful hosts for phosphorescent emitters.

Useful fluorescent dopants include, but are not limited to, derivatives of anthracene, tetracene, xanthene, perylene, rubrene, coumarin, rhodamine, and quinacridone, dicyanomethylenepyrans compounds, thiopyran compounds, polymethine compounds, pyrilium and thiapyrilium compounds,
20 fluorene derivatives, perflanthene derivatives, indenoperylene derivatives, bis(azinyl)amine boron compounds, bis(azinyl)methane compounds, and carbostyryl compounds.

Electron-Transporting Layer (ETL)

Preferred thin film-forming materials for use in forming the
25 electron-transporting layer 111 of the organic EL elements of this invention are metal chelated oxinoid compounds, including chelates of oxine itself (also commonly referred to as 8-quinolinol or 8-hydroxyquinoline). Such compounds help to inject and transport electrons, exhibit high levels of performance, and are readily fabricated in the form of thin films. Exemplary oxinoid compounds were
30 listed previously.

Other electron-transporting materials include various butadiene derivatives as disclosed in US 4,356,429 and various heterocyclic optical

brighteners as described in US 4,539,507. Benzazoles and triazines are also useful electron-transporting materials.

Cathode

When light emission is viewed solely through the anode, the cathode 113 used in this invention can be comprised of nearly any conductive material. Desirable materials have good film-forming properties to ensure good contact with the underlying organic layer, promote electron injection at low voltage, and have good stability. Useful cathode materials often contain a low work function metal (< 4.0 eV) or metal alloy. One preferred cathode material is comprised of a Mg:Ag alloy wherein the percentage of silver is in the range of 1 to 20 %, as described in US 4,885,221. Another suitable class of cathode materials includes bilayers comprising a thin electron-injection layer (EIL) in contact with the organic layer (e.g., ETL) which is capped with a thicker layer of a conductive metal. Here, the EIL preferably includes a low work function metal or metal salt, and if so, the thicker capping layer does not need to have a low work function. One such cathode is comprised of a thin layer of LiF followed by a thicker layer of Al as described in US 5,677,572. Other useful cathode material sets include, but are not limited to, those disclosed in US 5,059,861; 5,059,862, and 6,140,763.

When light emission is viewed through the cathode, the cathode must be transparent or nearly transparent. For such applications, metals must be thin or one must use transparent conductive oxides, or a combination of these materials. Optically transparent cathodes have been described in more detail in US 4,885,211, US 5,247,190, JP 3,234,963, US 5,703,436, US 5,608,287, US 5,837,391, US 5,677,572, US 5,776,622, US 5,776,623, US 5,714,838, US 5,969,474, US 5,739,545, US 5,981,306, US 6,137,223, US 6,140,763, US 6,172,459, EP 1 076 368, US 6,278,236, and US 6,284,393. Cathode materials are typically deposited by evaporation, sputtering, or chemical vapor deposition. When needed, patterning can be achieved through many well known methods including, but not limited to, through-mask deposition, integral shadow masking, for example, as described in US 5,276,380 and EP 0 732 868, laser ablation, and selective chemical vapor deposition.

Other Common Organic Layers and Device Architecture

In some instances, layers 109 and 111 can optionally be collapsed into a single layer that serves the function of supporting both light emission and electron transportation. It also known in the art that emitting dopants may be
5 added to the hole-transporting layer, which may serve as a host. Multiple dopants may be added to one or more layers in order to create a white-emitting OLED, for example, by combining blue- and yellow-emitting materials, cyan- and red-emitting materials, or red-, green-, and blue-emitting materials. White-emitting devices are described, for example, in EP 1 187 235, US 20020025419, EP 1 182
10 244, US 5,683,823, US 5,503,910, US 5,405,709, and US 5,283,182.

Additional layers such as electron or hole-blocking layers as taught in the art may be employed in devices of this invention. Hole-blocking layers are commonly used to improve efficiency of phosphorescent emitter devices, for example, as in US 20020015859.

15 This invention may be used in so-called stacked device architecture, for example, as taught in US 5,703,436 and US 6,337,492.

Deposition of organic layers

The organic materials mentioned above are suitably deposited through a vapor-phase method such as sublimation, but can be deposited from a
20 fluid, for example, from a solvent with an optional binder to improve film formation. If the material is a polymer, solvent deposition is useful but other methods can be used, such as sputtering or thermal transfer from a donor sheet. The material to be deposited by sublimation can be vaporized from a sublimator “boat” often comprised of a tantalum material, e.g., as described in US 6,237,529,
25 or can be first coated onto a donor sheet and then sublimed in closer proximity to the substrate. Layers with a mixture of materials can utilize separate sublimator boats or the materials can be pre-mixed and coated from a single boat or donor sheet. Patterned deposition can be achieved using shadow masks, integral shadow masks (US 5,294,870), spatially-defined thermal dye transfer from a donor sheet
30 (US 5,688,551, 5,851,709 and 6,066,357) and inkjet method (US 6,066,357).

Encapsulation

Most OLED devices are sensitive to moisture or oxygen, or both, so they are commonly sealed in an inert atmosphere such as nitrogen or argon, along with a desiccant such as alumina, bauxite, calcium sulfate, clays, silica gel, zeolites, alkaline metal oxides, alkaline earth metal oxides, sulfates, or metal halides and perchlorates. Methods for encapsulation and desiccation include, but are not limited to, those described in US 6,226,890. In addition, barrier layers such as SiO_x, Teflon, and alternating inorganic/polymeric layers are known in the art for encapsulation.

OLED devices of this invention can employ various well-known optical effects in order to enhance its properties if desired. This includes optimizing layer thicknesses to yield maximum light transmission, providing dielectric mirror structures, replacing reflective electrodes with light-absorbing electrodes, providing anti glare or anti-reflection coatings over the display, or providing colored, neutral density, or color conversion filters over the display. Filters, and anti-glare or anti-reflection coatings may be specifically provided over the encapsulating cover or an electrode protection layer beneath the encapsulating cover.

The invention has been described in detail with particular reference to certain preferred embodiments thereof, but it will be understood that variations and modifications can be effected within the spirit and scope of the invention.

PARTS LIST

10	OLED light emissive element
11	top-emitting OLED display device
12	organic light emitting layer
12R	red-light emitting OLED layer
12G	green-light emitting OLED layer
12B	blue-light emitting OLED layer
14	second electrode layer
16	first electrode layer
18	power source
20	substrate
22	TFT active matrix layer
24	first insulating planarization layer
24'	second insulating planarization layer
26	hole-injecting layer
27	hole-transporting layer
28	electron-transporting layer
29	electron-injection layer
32	electrode-protection layer
34	cavity
36	encapsulating cover
37	anti-reflection coating
38	optional protective cover
50	circular light polarizer
52	peripheral channel
70	adhesive
101	substrate
103	anode layer
105	hole-injecting layer
107	hole-transporting layer
109	light-emitting layer
111	electron-transporting layer
113	cathode layer
250	voltage/current source
260	conductive wiring